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=> s NTP1 or neuronal tyrosine/threonine phosphatase
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=> s NTP1 or neuronal tyrosine/threonine phosphatase
? TRUNCATION SYMBOL NOT VALID WITHIN 'TYROSINE?THREONINE'
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? TRUNCATION SYMBOL NOT VALID WITHIN 'TYROSINE?THREONINE'
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=> s NTP1 or neuronal tyrosine/threonine phosphatase
L1 2 NTP1 OR NEURONAL TYROSINE/THREONINE PHOSPHATASE

=> s ! and (transgen? or knockout or disrupt? or deficient? or delet?)
L2 1 L1 AND (TRANSGEN? OR KNOCKOUT OR DISRUPT? OR
DEFICIENT? OR DELET?)
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L2 ANSWER 1 OF 1 CAPLUS COPYRIGHT 2003 ACS
AN 2002 638352 CAPLUS
DN 137 180791
TI ***Transgenic*** mice containing neuronal tyrosine/threonine protein
phosphatase gene ***NTP1*** and their use as
disease models and for screening for modulators
IN Allen, Keith D
PA USA
SO U.S. Pat. Appl. Publ., 26 pp
CODEN USXXCO
DT Patent
LA English
FAN CNT 1

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI US 2002116729	A1	20020822	US 2001-5858	20011204
PRAI US 2000-251802P	P	20001206		
AB The present invention relates to ***transgenic*** animals, as well as comprns. and methods relating to the characterization of gene function. Specifically, the present invention provides ***transgenic*** mice comprising a ***disruption*** in the ***NTP1*** gene encoding a neuronal tyrosine/threonine phosphatase, a member of the mitogen-activated protein kinase phosphatase gene family which contains a complex trinucleotide repeat in the coding region. To investigate the role of ***NTP1***, ***disruptions*** in the ***NTP1*** genes are produced by homologous recombination using 5' and 3' arms in a targeting construct ***Transgenic*** mice contg. ***NTP1*** ***disruptions*** exhibit anti-depressive behavior, relative to wild type mice, as shown by a decrease in immobile time when tail suspended. Such ***transgenic*** mice are useful as models for disease and for identifying agents that modulate gene expression and gene function, and as potential treatments for various disease states and disease conditions.				

=> d bib abs 11

L1 ANSWER 1 OF 2 CAPLUS COPYRIGHT 2003 ACS
AN 2002 638352 CAPLUS
DN 137 180791
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gene ***NTP1*** disruptions and their use as disease models and for
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L1 ANSWER 1 OF 2 CAPLUS COPYRIGHT 2003 ACS

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L1 ANSWER 2 OF 2 CAPLUS COPYRIGHT 2003 ACS

AN 2001 447761 CAPLUS

DN 135 180233

TI The in vivo neuromodulatory effects of the herbal medicine Ginkgo biloba
AU Watanabe, Coran M. H.; Wolffram, Siegfried, Ader, Peter; Rimbach, Gerald; Packer, Lester; Maguire, John J.; Schultz, Peter G.; Gohil, Kishorchandra
CS Department of Chemistry, Scripps Research Institute, La Jolla, CA, 92037, USA

SO Proceedings of the National Academy of Sciences of the United States of America (2001), 98(12), 6577-6580

CODEN PNASA6, ISSN 0027-8424

PB National Academy of Sciences

DT Journal

LA English

AB Exts of G biloba leaves are consumed as dietary supplements to counteract chronic age-related neuro disorders. High-d oligonucleotide microarrays were used to define the transcriptional effects in the brain cortex and hippocampus of adult female C57BL/6 mice fed diets supplemented with the herbal ext. Gene expression RT-PCR anal. was then focused on the mRNAs that showed >3-fold change in their expression. In the brain cortex, mRNAs for neuronal tyrosine/threonine phosphatase 1 and microtubule-assocd. protein factor tau were enhanced. Hyperphosphorylated tau is the major constituent of the neurofibrillary tangles in the brain of Alzheimer disease patients. The expression of alpha-amino-3-hydroxy-5-methyl-4-isoxazolepropionic acid channels (AMPA-2 receptor), calcium and chloride channels, prolactin, and growth hormone (GH), all of which are assocd with brain function, were also up-regulated. In the hippocampus, only transthyretin mRNA was upregulated. Transthyretin has a role in hormone transport in the brain and possibly a neuroprotective role by amyloid-beta sequestration. Thus, diets supplemented with G biloba leaf ext. have notable neuromodulatory effects in vivo. The data illustrate the utility of genome-wide expression monitoring to investigate the biol. actions of complex herbal exts.

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ALL CITATIONS AVAILABLE IN THE RE FORMAT

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